

Original citation:

Pöppler, Ann Christin, Corletta, Emily K., Pearce, Harriet, Seymour, Mark, Reid, Matthew M., Montgomery, Mark G. and Brown, Steven P.. (2017) Single-crystal X-ray diffraction and NMR crystallography of a 1:1 cocrystal of dithianon and pyrimethanil. Acta Crystallographica Section C : Structural Chemistry, C73 . pp. 149-156.

Permanent WRAP URL:

<http://wrap.warwick.ac.uk/85429>

Copyright and reuse:

The Warwick Research Archive Portal (WRAP) makes this work of researchers of the University of Warwick available open access under the following conditions.

This article is made available under the Creative Commons Attribution 2.0 Generic (CC BY 2.0) license and may be reused according to the conditions of the license. For more details see: <http://creativecommons.org/licenses/by/2.0/>

A note on versions:

The version presented in WRAP is the published version, or, version of record, and may be cited as it appears here.

For more information, please contact the WRAP Team at: wrap@warwick.ac.uk

Single-crystal X-ray diffraction and NMR crystallography of a 1:1 cocrystal of dithianon and pyrimethanil

Ann-Christin Pöppler,^{a,b} Emily K. Corlett,^{a,c} Harriet Pearce,^{a,c} Mark P. Seymour,^d Matthew Reid,^{d,e} Mark G. Montgomery^d and Steven P. Brown^{a*}

Received 30 September 2016

Accepted 17 January 2017

Edited by D. L. Bryce, University of Ottawa, Canada

Keywords: NMR crystallography; solid-state NMR; dithianon; pyrimethanil; cocrystal; hydrogen bonding; C—H... π interactions; fungicides.

CCDC reference: 1507863

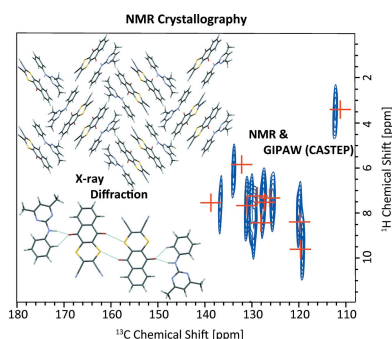
Supporting information: this article has supporting information at journals.iucr.org/c

^aDepartment of Physics, University of Warwick, Coventry CV4 7AL, United Kingdom, ^bDepartment of Organic Chemistry, University of Würzburg, 97074 Würzburg, Germany, ^cMolecular Analytical Science Centre for Doctoral Training, University of Warwick, Coventry CV4 7AL, United Kingdom, ^dInternational Research Centre, Syngenta, Jealott's Hill, Bracknell, Berkshire RG42 6EY, United Kingdom, and ^eAfton Chemical, London Road, Bracknell, Berkshire RG12 2UW, United Kingdom. *Correspondence e-mail: s.p.brown@warwick.ac.uk

A single-crystal X-ray diffraction structure of a 1:1 cocrystal of two fungicides, namely dithianon (DI) and pyrimethanil (PM), is reported [systematic name: 5,10-dioxo-5*H*,10*H*-naphtho[2,3-*b*][1,4]dithiine-2,3-dicarbonitrile-4,6-dimethyl-*N*-phenylpyrimidin-2-amine (1/1), C₁₄H₄N₂O₂S₂·C₁₂H₁₃N₂]. Following an NMR crystallography approach, experimental solid-state magic angle spinning (MAS) NMR spectra are presented together with GIPAW (gauge-including projector augmented wave) calculations of NMR chemical shieldings. Specifically, experimental ¹H and ¹³C chemical shifts are determined from two-dimensional ¹H–¹³C MAS NMR correlation spectra recorded with short and longer contact times so as to probe one-bond C–H connectivities and longer-range C...H proximities, whereas H...H proximities are identified in a ¹H double-quantum (DQ) MAS NMR spectrum. The performing of separate GIPAW calculations for the full periodic crystal structure and for isolated molecules allows the determination of the change in chemical shift upon going from an isolated molecule to the full crystal structure. For the ¹H NMR chemical shifts, changes of 3.6 and 2.0 ppm correspond to intermolecular N–H...O and C–H...O hydrogen bonding, while changes of –2.7 and –1.5 ppm are due to ring current effects associated with C–H... π interactions. Even though there is a close intermolecular S...O distance of 3.10 Å, it is of note that the molecule-to-crystal chemical shifts for the involved sulfur or oxygen nuclei are small.

1. Introduction

With an increasing global population, limited availability of arable land, an increase in extreme weather events and growing pest resistance to certain existing agrochemical products, innovation in the agrochemical industry is as important as ever if we are to provide enough food for everyone. With lower usage rates, ease of use and more favourable toxicology profiles being important objectives, the search for and structure-based design of potential agrochemical products needs to become more efficient (Lamberth *et al.*, 2013). One possibility in this regard is the usage of cocrystals formed between an active ingredient and coformers or other active ingredients *via* reversible noncovalent interactions. While this is an established procedure in the development of new active pharmaceutical ingredients, where it is used to increase the solubility and bioavailability (Blagden *et al.*, 2007), there is also great potential to exploit cocrystals in the optimization and development of agrochemicals. For example, a reduced solubility could increase the agrochemical's resi-



OPEN ACCESS

Table 1
Experimental details.

Crystal data	
Chemical formula	$C_{14}H_4N_2O_2S_2 \cdot C_{12}H_{13}N_3$
M_r	495.59
Crystal system, space group	Monoclinic, $P2_1/n$
Temperature (K)	100
a, b, c (Å)	7.1707 (2), 22.8006 (6), 13.8237 (4)
β (°)	97.047 (3)
V (Å ³)	2243.04 (7)
Z	4
Radiation type	Cu $K\alpha$
μ (mm ⁻¹)	2.45
Crystal size (mm)	0.60 × 0.10 × 0.02
Data collection	
Diffractometer	Agilent Xcalibur Onyx Ultra
Absorption correction	Multi-scan (CrysAlis PRO; Agilent, 2014)
T_{min}, T_{max}	0.596, 1.000
No. of measured, independent and observed [$I > 2.0\sigma(I)$] reflections	5143, 3160, 2667
R_{int}	0.035
θ_{max} (°)	58.9
$(\sin \theta/\lambda)_{max}$ (Å ⁻¹)	0.556
Refinement	
$R[F^2 > 2\sigma(F^2)], wR(F^2), S$	0.045, 0.094, 0.98
No. of reflections	3141
No. of parameters	109
No. of restraints	3
H-atom treatment	H atoms treated by a mixture of independent and constrained refinement
$\Delta\rho_{max}, \Delta\rho_{min}$ (e Å ⁻³)	0.43, -0.37

Computer programs: CrysAlis PRO (Agilent, 2014), SUPERFLIP (Palatinus & Chapuis, 2007), CRYSTALS (Betteridge *et al.*, 2003), CAMERON (Watkin *et al.*, 1996) and Mercury (Macrae *et al.*, 2006).

dence time on the respective plant and multicomponent entities could improve the release profile (and thus absolute usage), as well as allow the simultaneous delivery of two or more active components. However, the design of suitable cocrystalline materials and prediction of their properties and formed cocrystal structures is far from being trivial. Some design strategies based on the hierarchy of intermolecular interactions (Aakeroy & Salmon, 2005) or the assessment of the solubilities and saturation temperatures of the pure compounds to be included in a cocrystalline arrangement (ter Horst *et al.*, 2009) are available as a guideline. However, if multiple and different hydrogen-bonding donors and acceptors are present in the molecules, a reliable prediction of the resulting structure becomes very difficult (Bhatt *et al.*, 2009).

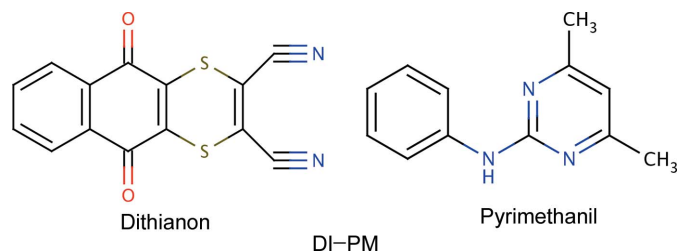
NMR crystallography, namely the combination of experimental solid-state magic angle spinning (MAS) NMR with calculation of NMR parameters, is finding important application to moderately sized organic molecules (Harris, 2004; Elena *et al.*, 2006; Harris *et al.*, 2009; Bonhomme *et al.*, 2012). We present here an NMR crystallography analysis of the 1:1 cocrystal of two fungicides, namely dithianon (DI) and pyrimethanil (PM). Specifically, following a preparation protocol in Sowa *et al.* (2013), a single-crystal X-ray diffraction structure determination is reported, with this structure (after DFT geometry optimization) providing the input for a calculation, using the GIPAW (gauge-including projector augmented

wave) method (Pickard & Mauri, 2001; Yates *et al.*, 2007), of the NMR chemical shieldings. The computational analysis is complemented by the recording of 1D (one-dimensional) and 2D (two-dimensional) experimental ¹H and ¹³C MAS NMR spectra. Building upon studies of pharmaceutical cocrystals by such an NMR crystallography investigation (Tatton *et al.*, 2013; Dudenko *et al.*, 2013; Stevens *et al.*, 2014; Kerr *et al.*, 2015; Sardo *et al.*, 2015; Luedeker *et al.*, 2016), we present here the application of this approach to an agrochemical cocrystal.

2. Experimental and computational details

2.1. Sample preparation

The DI–PM cocrystal was prepared according to method VII in point [0041] of Sowa *et al.* (2013), *i.e.* dry dithianon and pyrimethanil (both solids) were mixed thoroughly in a 1:1 molar ratio (0.5 g of pyrimethanil) and kept at 323 K under agitation. After a couple of hours, the powdery product had changed to a dark-olive-green colour.



2.2. Single-crystal X-ray diffraction: structure solution and refinement

Crystal data, data collection and structure refinement details are summarized in Table 1. The H atoms were all located in a difference map, but those attached to C atoms were repositioned geometrically. The H atoms were initially refined with soft restraints on the bond lengths and angles to regularize their geometry [$C-H = 0.93-0.98$ Å and $N-H = 0.86-0.89$ Å, and with $U_{iso}(H) = 1.2-1.5U_{eq}(\text{parent})$], after which the positions were refined with riding constraints (Cooper *et al.*, 2010).

2.3. Solid-state NMR

1D ¹H MAS and 1D ¹³C cross polarization (CP) MAS experiments were performed on a Bruker Avance III spectrometer operating at ¹H and ¹³C Larmor frequencies of 600 and 150.9 MHz, respectively, using a 1.3 mm HXY (¹H MAS) or a 4 mm HX (¹³C CP MAS) Bruker probe. In all cases, a ¹H 90° pulse duration of 2.5 µs was used. 2D ¹H–¹³C HETCOR experiments were performed on a Bruker Avance III spectrometer, using a 4 mm HXY probe in double-resonance mode. In the HETCOR pulse sequence, the following phase cycling was employed: ¹H 90° pulse (90° 270°), ¹³C 180° pulse (2{0°} 2{180°}), ¹³C CP contact pulse (4{0°} 4{180°} 4{90°} 4{270°}), receiver (0° 180° 0° 180° 180° 0° 180° 0° 90° 270° 90° 270° 270° 90° 270° 90°). For CP, a 70 to 100% ramp (Metz *et al.*, 1994) on the ¹H channel was used for the CP contact time. During acquisition of a ¹³C FID, SPINAL64 (Fung *et al.*, 2000)

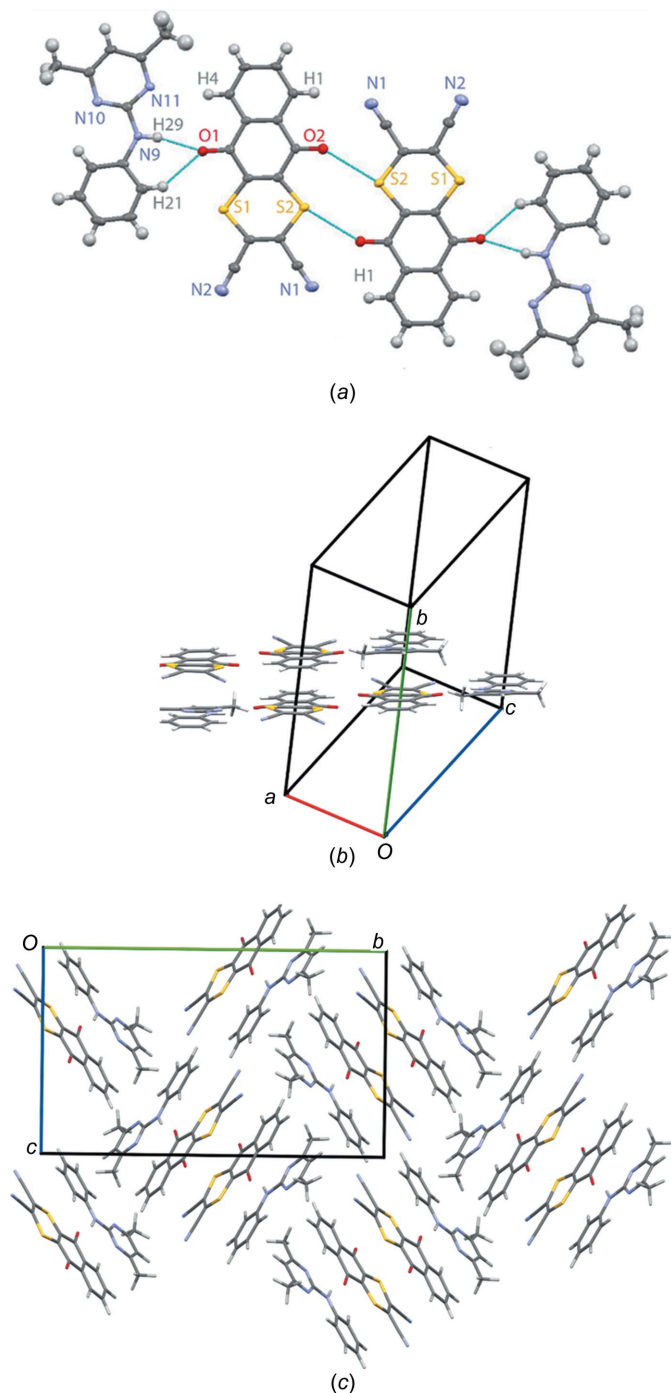


Figure 1

Representations of the crystal structure of the DI-PM cocrystal, showing (a) the intermolecular interactions within a 'chain' of molecules, with displacement ellipsoids drawn at the 50% probability level, (b) the packing of two chains of molecules as 'layers' and (c) the 'zigzag' arrangement of chains (viewed along the crystallographic *a* axis). In parts (b) and (c), the unit cell is shown, indicating the *a*, *b* and *c* unit-cell axes.

^1H heteronuclear decoupling was applied with a pulse duration of 5.9 μs at a nutation frequency of 100 kHz. A 2D ^1H DQ experiment with BABA recoupling (Sommer *et al.*, 1995; Schnell *et al.*, 1998) was performed on a Bruker Avance III spectrometer operating at a ^1H Larmor frequency of 700 MHz using a 1.3 mm HXY Bruker probe. A 16-step phase cycle was

used to select $\Delta p = \pm 2$ on the DQ excitation block and $\Delta p = -1$ on the z -filter 90° pulse, where p is the coherence order. In all 2D experiments, the States-TPPI method was used to achieve sign discrimination in F_1 . ^{13}C and ^1H chemical shifts are referenced with respect to TMS using L-alanine at natural abundance as an external reference: 177.8 ppm for the ^{13}C carboxylate resonance and 1.1 ppm for the ^1H methyl resonance. All experiments were performed at room temperature, though frictional effects due to MAS increase the actual sample temperature (Langer *et al.*, 1999).

2.4. DFT calculations

Calculations were performed using CASTEP (Clark *et al.*, 2005; Academic Release Version 8.0) and employed the PBE exchange-correlational functional (Perdew *et al.*, 1996). For both geometry optimization and NMR shielding calculations, a plane-wave basis set with ultrasoft pseudopotentials (Vanderbilt, 1990) with a maximum plane-wave cut-off energy of 700 eV was used. A Monkhorst-Pack grid of minimum sample spacing $0.05 \times 2\pi \text{ \AA}^{-1}$ was used to take integrals over the Brillouin zone. Geometry optimization was performed with the unit-cell parameters fixed, starting from the single-crystal X-ray structure. The positions of the 208 atoms in the unit cell ($Z = 4$, $Z' = 1$) were relaxed and periodic boundary conditions were applied. The space group $P2_1/n$ was preserved. All distances and angles stated in the main text of this article are for the geometry-optimized crystal structure. Note also that the geometry optimization within CASTEP causes a relabelling of the atoms – in this article, we use the CASTEP numbering; see Fig. S1 in the *Supporting information* for a comparison with the numbering employed in the crystallographic CIF file. The GIPAW method (Pickard & Mauri, 2001; Yates *et al.*, 2007) was utilized for the NMR chemical-shielding calculations, which were performed on the geometry-optimized structure. For the isolated molecule calculations, a single molecule (either DI or PM) from the fully geometry optimized structure is kept in the unit cell, whose dimensions are also increased by $\sim 5 \text{ \AA}$ in each direction – the NMR shieldings are then calculated without any further geometry optimization.

3. Results and discussion

3.1. Single-crystal X-ray diffraction structure

The single-crystal X-ray diffraction structure of the DI-PM cocrystal is schematically represented in Fig. 1. As shown in Fig. 1(a), a chain of molecules is held together by $\text{N}-\text{H} \cdots \text{O}$ and $\text{C}-\text{H} \cdots \text{O}$ hydrogen bonds (between DI and PM molecules) and by putative $\text{S} \cdots \text{O}$ interactions (Burling & Goldstein, 1992) between two DI molecules; note that the relative strengths of these interactions is investigated below (see §3.5) using GIPAW calculations of NMR chemical shieldings. The further packing of two chains of molecules as 'layers' and a 'zigzag' arrangement of chains are shown in Figs. 1(b) and 1(c), respectively. As can be seen from the representation along the crystallographic *a* axis in Fig. 1(c), the packing is

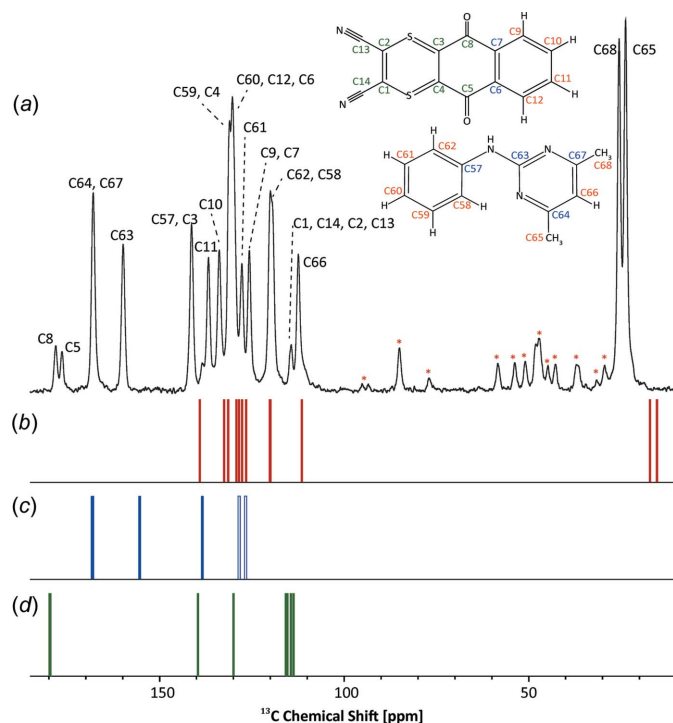


Figure 2
(a) A ^1H (600 MHz)- ^{13}C CP MAS (12.5 kHz) NMR spectrum of the DI-PM cocrystal (* denote spinning sidebands), together with (b)–(d) stick spectra corresponding to calculated (GIPAW) ^{13}C chemical shifts (see Table 2). Separate stick spectra are presented according to whether correlation peaks corresponding to (b) direct C–H bonds or (c) longer-range C...H proximities are observed in the ^1H - ^{13}C 2D spectra presented in Fig. 4, or (d) where no experimental correlation peaks are observed. In the CP MAS experiment, a contact time of 1.4 ms was used and 1024 transients were co-added for a recycle delay of 57 s.

based on assemblies of blocks of four molecules; four molecules (PM-DI-DI-PM) are arranged in a layer (Fig. 1a), forming a block that is perpendicular to an adjacent block of four molecules, thus building up the ‘zigzag’ arrangement.

3.2. Experimental and calculated ^{13}C chemical shifts

Fig. 2 presents a ^{13}C CP MAS NMR 1D spectrum (Fig. 2a) of the DI-PM cocrystal, together with three stick spectra (Figs. 2b, 2c and 2d) that represent ^{13}C chemical shifts calculated using the GIPAW method for the DI-PM crystal structure. Specifically, the calculated ^{13}C chemical shifts are presented in three groups according to whether they correspond to direct one-bond C–H connectivities (Fig. 2b, red labels) or nonprotonated C atoms (Figs. 2c and 2d, blue and green labels, respectively). The distinction between Figs. 2(c) and 2(d) corresponds to whether cross peaks corresponding to a longer-range C...H proximity are observed in ^1H - ^{13}C 2D correlation spectra (see §3.4).

3.3. One- and two-dimensional ^1H MAS NMR spectra

Figs. 3(a) and 3(b) present ^1H NMR spectra of the DI-PM cocrystal recorded at a fast MAS frequency of 60 kHz; specifically, a one-pulse one-dimensional spectrum in Fig. 3(a), together with vertical lines corresponding to calculated

Table 2

Comparison of calculated (GIPAW)^a and experimental ^{13}C and ^1H NMR chemical shifts (in ppm) in the DI-PM cocrystal^b.

Atom label		^{13}C		^1H	
C	H	δ_{calc}	δ_{expt}	δ_{calc}	δ_{expt}
C65	H22/H23/H24 ^c	15.3	23.9	1.8	1.9
C68	H26/H27/H28 ^c	17.2	25.7	2.0	2.0
C66	H25	111.5	112.6	3.4	4.0
C1	–	113.8	114.4 ^d	–	–
C14	–	114.5	114.4 ^d	–	–
C2	–	115.5	114.4 ^d	–	–
C13	–	115.9	114.4 ^d	–	–
C58	H17	120.1	119.4	9.7	9.1
C62	H21	120.2	120.3	8.4	8.0
C9	H1	126.7	125.7	7.4	7.4
C7	H1 ^e	126.8	125.7	7.4	7.4
C61	H20	127.7	127.7	7.6	7.4
C12	H4	128.5	129.8	8.5	8.2
C6	H4 ^e	128.6	129.8	8.5	8.2
C60	H19	129.3	130.2	7.3	7.8
C4	–	130.1	131.1 ^d	–	–
C59	H18	131.5	131.2	7.7	7.7
C10	H2	132.6	133.9	5.9	6.2
C11	H3	139.2	136.8	7.6	7.7
C57	H21, H17, H29	138.5	141.5	8.4, 9.7, 10.5	8.9
C3	–	139.7	141.4 ^d	–	–
C63	H29	155.5	160.1	10.5	9.1
C67	H26/H27/H28, H25	168.2	168.2	2.0, 3.4	2.8
C64	H22/H23/H24, H25	168.4	168.2	1.8, 3.4	2.8
C5	–	179.7	176.5 ^d	–	–
C8	–	179.9	178.2 ^d	–	–

Notes: (a) calculated isotropic chemical shifts are determined from calculated chemical shieldings according to $\delta_{\text{calc}} = \sigma_{\text{ref}} - \sigma_{\text{calc}}$, where σ_{ref} equals 30.0 ppm for ^1H and 163.2 ppm for ^{13}C . (b) H-atom labels and calculated and experimental ^1H chemical shifts are presented in normal font for direct one-bond C–H connectivities, while longer-range C...H proximities (corresponding to cross peaks observed in the ^1H - ^{13}C spectra presented in Figs. 4b and 4c) are presented in italics. (c) For CH_3 groups, the calculated ^1H chemical shifts correspond to the average over the three H atoms. (d) Experimental chemical shifts taken from the ^{13}C CP MAS spectrum (Fig. 2a) since no cross peaks are observed in the ^1H - ^{13}C spectra presented in Figs. 4(b) and 4(c). (e) Note that the C7–H1 and C6–H4 cross peaks due to longer-range C...H proximities cannot be distinguished from the C9–H1 and C12–H4 cross peaks due to one-bond C–H connectivities – in the stick spectrum in Fig. 2(c), open bars denote the calculated (GIPAW) C7 and C6 ^{13}C chemical shifts.

(GIPAW) ^1H chemical shifts, as well as a 2D DQ spectrum in Fig. 3(b). In addition, Fig. 3(c) presents a ^1H - ^{13}C 2D correlation spectrum of the DI-PM cocrystal; note that this spectrum has been rotated through 90° from its usual representation such that the direct (^{13}C) dimension is vertical. In this way, it is possible to directly compare (see vertical dashed lines) ^1H chemical shifts of peaks in the ^1H - ^{13}C (Fig. 3c) and ^1H DQ 2D (Fig. 3b) and ^1H 1D (Fig. 3a) spectra. Two separate spectral regions are presented in Fig. 3(c) corresponding to (top) the methyl resonances at a ^{13}C chemical shift close to 25 ppm and (bottom) the aromatic CH resonances with ^{13}C chemical shifts between 110 and 140 ppm.

The ^1H - ^{13}C correlation spectrum in Fig. 3(c) was recorded using a short CP contact time of 100 μs to transfer magnetization from ^1H to ^{13}C , such that cross peaks correspond to one-bond C–H connectivities. The spreading of the resonances into two dimensions in Fig. 3(c) allows the identification of two and ten resolved cross peaks for the CH_3 and aromatic CH groups, respectively. The value of such a ^1H - ^{13}C correlation spectrum in resolving and assigning the experimental ^1H chemical shifts is thus evident. Table 2 lists the calculated (GIPAW) and experimental ^{13}C chemical shifts

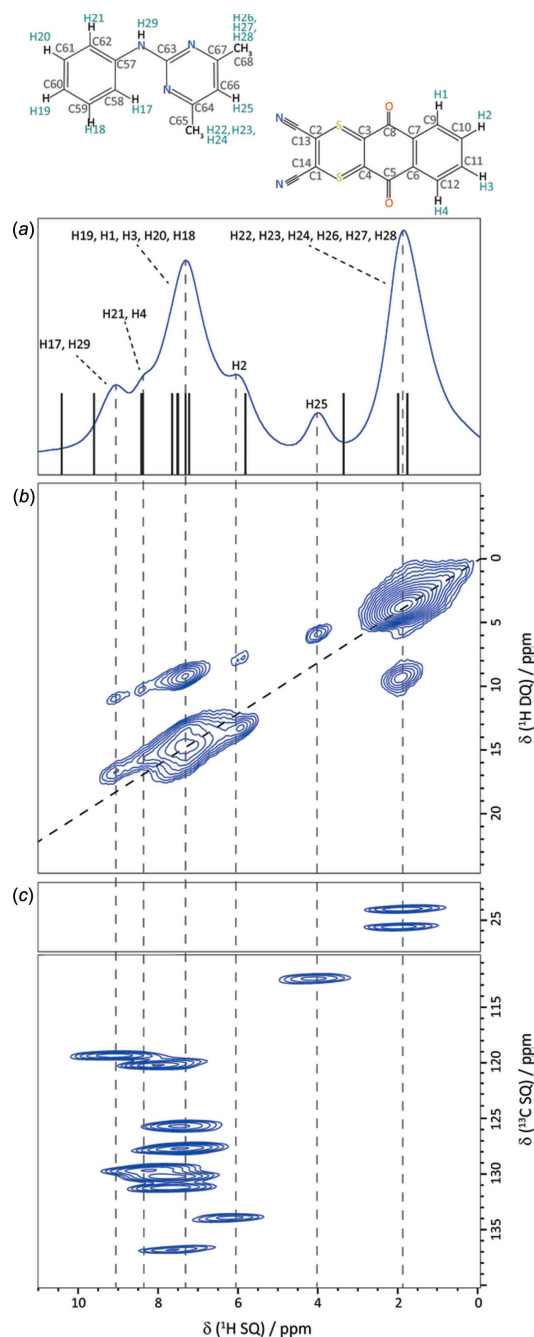


Figure 3

MAS NMR spectra of the DI-PM cocrystal, showing (a) a ^1H (600 MHz) MAS (60 kHz) one-pulse spectrum (16 transients were co-added for a recycle delay of 15 s), (b) a 2D ^1H (700 MHz) DQ MAS (60 kHz) spectrum (the dashed diagonal line indicates the $F_1 = 2F_2$ DQ-SQ diagonal) recorded using one rotor period of BABA recoupling (32 transients were co-added for each of 200 t_1 FIDs using a recycle delay of 6 s, corresponding to a total experiment time of 12 h) and (c) a ^1H (500 MHz)- ^{13}C HETCOR MAS (12.5 kHz) spectrum recorded using FSLG ^1H homonuclear decoupling in t_1 and a short CP transfer duration of 100 μs (104 transients were co-added for each of 128 t_1 FIDs using a recycle delay of 6 s, corresponding to a total experimental time of 22 h). The vertical lines in part (a) correspond to calculated (GIPAW) ^1H chemical shifts. For the ^1H - ^{13}C NMR spectrum in part (c), two separate spectral regions are presented corresponding to methyl and aromatic C-H groups; note that this spectrum has been rotated through 90° from its usual representation [the ^{13}C dimension corresponds to direct (t_2) acquisition]. The base contour level is at (b) 7% and (c) 20% of the maximum peak height.

(sorted in order of increasing chemical shift). For directly bonded C-H connectivities, H-atom labels and calculated (GIPAW) and experimental ^1H chemical shifts are presented in normal font.

Fig. 4 compares ^1H - ^{13}C correlation spectra recorded with three different CP contact times of 100 μs (Fig. 4a), 500 μs (Fig. 4b) and 1 ms (Fig. 4c); Fig. 4(a) is a copy of Fig. 3(c), but presented in the normal orientation, *i.e.* with the direct (^{13}C) dimension horizontal. It is evident that additional cross peaks are observed for longer CP contact times – these correspond to longer-range C...H proximities (see italics font in Table 2). Notably, cross peaks are observed at ^{13}C chemical shifts of 141.5 (atom C57), 160.1 (atom C63) and 168.2 ppm (atoms C64 and C67); these all correspond to intramolecular proximities within the dithianon molecule, *i.e.* C57 with H17 (9.1 ppm, 2.16 Å), H21 (8.0 ppm, 2.16 Å) and H29 (9.1 ppm, 2.06 Å), C63 with H29 (9.1 ppm, 2.01 Å), C64 and C67 with H25 (4.0 ppm, 2.16 and 2.17 Å) and CH_3 protons (1.9 and 2.0 ppm, nearest distance 2.14 Å). Of most interest is the (160.1 ppm, 9.1 ppm) cross peak, which thus enables the determination of the NH ^1H chemical shift.

With all the ^1H chemical shifts assigned, let us re-examine the ^1H DQ MAS spectrum in Fig. 3(b). In such a spectrum, cross peaks are observed in the DQ dimension at the sum of the two single-quantum (SQ) frequencies if there is a close proximity (typically up to 3.5 Å; Brown, 2007, 2012) between the corresponding two H atoms (a full listing of H...H proximities under 3.5 Å for the DI-PM cocrystal is given in Table S1 of the *Supporting information*). Consider the two lowest-ppm aromatic CH protons H25 (4.0 ppm) and H2 (6.2 ppm) for which distinct ^1H resonances are resolved in the ^1H SQ dimension. For H25, the only DQ peak is at $4.0 + 2.0 = 6.0$ ppm with the CH_3 protons, since H25 is sandwiched between two methyl-group substituents on the PM molecule. For H2, there is a DQ peak at $6.2 + 7.5 = 13.7$ ppm corresponding to the intramolecular H...H proximity with the neighbouring H1 (7.4 ppm, 2.50 Å) and H3 (7.7 ppm, 2.47 Å) DI aromatic CH protons, as well as a DQ peak at $6.2 + 2.0 = 8.2$ ppm due to intermolecular proximities to the PM CH_3 H atoms (H23, H24, H28 and H22 at 2.90, 3.03, 3.12 and 3.12 Å, respectively). Considering the high-ppm region, DQ cross peaks for the overlapping PI NH H29 (9.1 ppm) and aromatic CH H17 (9.1 ppm) resonances are observed at $9.1 + 7.7 = 16.8$ ppm for intramolecular H29...H21 (2.21 Å) and H17...H18 (2.50 Å) proximities, as well as at $9.1 + 2.0 = 11.1$ ppm for intermolecular proximities to PM methyl-group protons (closest distances of H17...H26 = 2.48 Å and H29...H24 = 2.64 Å). For the other overlapping CH aromatic resonances, cross peaks due to intramolecular proximities with other CH aromatic resonances, as well as intermolecular proximities to the methyl protons, are also observed.

3.4. Comparison of experimental and calculated ^1H and ^{13}C chemical shifts

In the ^1H - ^{13}C correlation spectra presented in Fig. 4, red crosses correspond to calculated (GIPAW) ^{13}C and ^1H

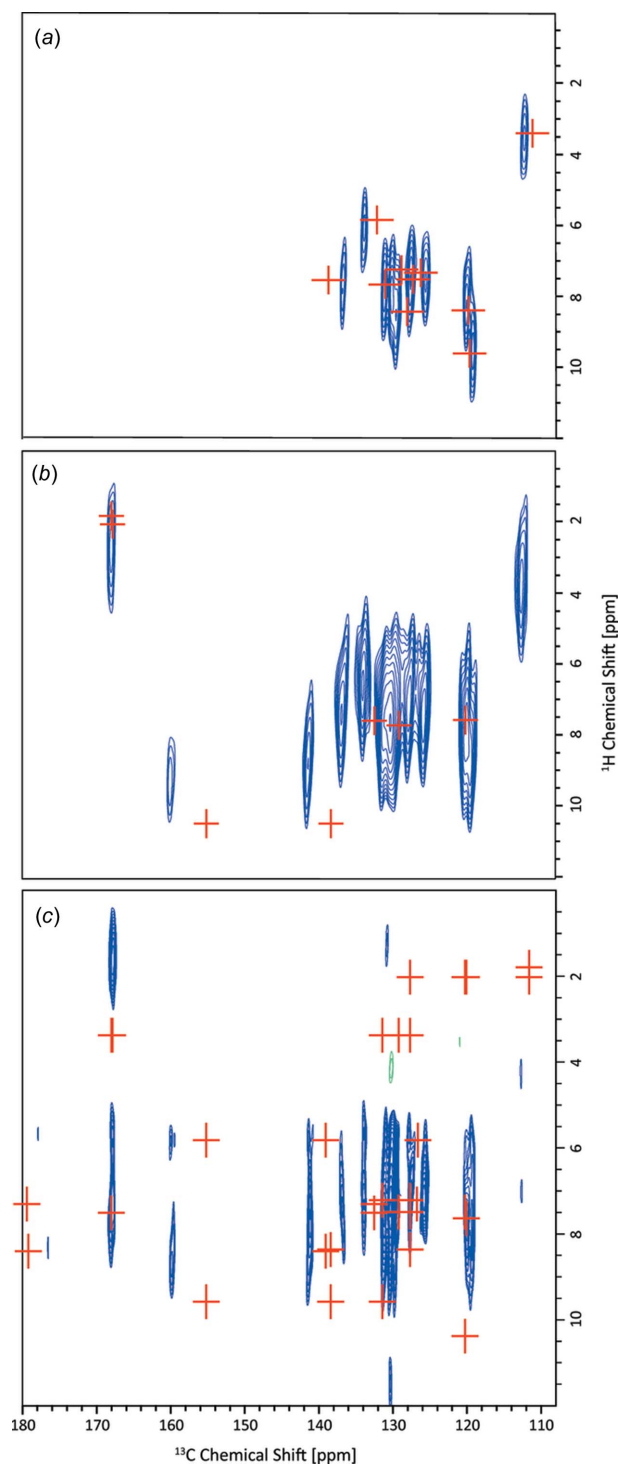


Figure 4
 ^1H (500 MHz)– ^{13}C HETCOR MAS (12.5 kHz) spectra of the DI–PM cocrystal recorded using FSLG ^1H homonuclear decoupling (Bielecki *et al.*, 1989) in t_1 with a CP transfer duration of (a) 100 μs , (b) 500 μs and (c) 1 ms. The spectrum in part (a) is repeated from Fig. 3(c). 104 transients were co-added for each of (b) 128 or (c) 90 t_1 FIDs using a recycle delay of (b) 6 or (c) 5.5 s, corresponding to a total experimental time of (b) 22 or (c) 14 h. The scaling factor in F_1 was determined to be (a) and (b) 1.80 or (c) 1.73. The base contour level is at (a) 20, (b) 13 and (c) 25% of the maximum peak height. Red crosses correspond to GIPAW-calculated ^1H and ^{13}C chemical shifts (see Table 2) for (a) one-bond C–H bonds and (b) and (c) C···H proximities between (b) 1.2 and 2.2 Å, and (c) 2.2 and 3.0 Å.

chemical shifts. Specifically, in Fig. 4(a), red crosses correspond to direct C–H one-bond connectivities (C–H distances under 1.2 Å), while in Figs. 4(b) and 4(c), red crosses are presented for C–H proximities between 1.2 and 2.2 Å (Fig. 4b), and between 2.2 and 3.0 Å (Fig. 4c). We comment here on the level of agreement between experimental and calculated (GIPAW) chemical shifts. Starting with a consideration of the aromatic CH moieties (see Fig. 4a and Table 2), the discrepancy between experiment and calculation is within 2 ppm for the ^{13}C chemical shifts (except for C11, where the difference is 2.4 ppm); this corresponds to the established observation that the discrepancy is within 1% of the chemical shift range (~ 200 ppm for ^{13}C chemical shifts of diamagnetic molecules). For the ^1H chemical shifts, while most are within the usual 0.3 ppm, some exhibit slightly larger discrepancies, notably 0.6 ppm for atoms H17 and H25.

For the two CH_3 groups (see Figs. 2 and 3a, and Table 2), there is excellent agreement for the ^1H chemical shifts (within 0.1 ppm), whereas the calculated ^{13}C chemical shifts are both 8.5 ppm lower than the experimental values, although the experimental difference in ^{13}C chemical shifts between atoms C65 and C68 of 1.8 ppm is reproduced by the calculation (difference of 1.9 ppm). The explanation for this is well known, namely, the gradient of a plot of experimental ^{13}C chemical shifts against calculated shielding deviates slightly from -1 (Harris *et al.*, 2007; Ashbrook & McKay, 2016), such that calculated ^{13}C chemical shifts are too low and too high compared to experiment for low-ppm and high-ppm resonances if, as here (see Fig. 2), the gradient is constrained to -1 and a single reference shielding is used. An alternative approach would be to use different reference shieldings for different regions of the spectrum (Webber, Emsley *et al.*, 2010).

Returning to the ^1H chemical shifts, the biggest discrepancy is for the NH proton (H29), where the calculated ^1H chemical shift of 10.5 ppm is 1.4 ppm higher than the experimental value of 9.1 ppm. Such a large difference is explained by a known temperature dependence (the experimental ^1H chemical shift increases upon reducing the temperature) for hydrogen-bonded protons (Brown *et al.*, 2001; Pickard *et al.*, 2007; Webber, Elena *et al.*, 2010), considering that the calculation corresponds to 0 K.

3.5. Calculated molecule-to-crystal changes in chemical shifts

For cases such as the DI–PM cocrystal in this article, an NMR crystallography study is able to provide new insight by means of a comparison of chemical shifts calculated for the full crystal structure with those calculated for an isolated molecule (as extracted from the geometry-optimized crystal structure) (Yates *et al.*, 2005; Schmidt *et al.*, 2006; Mafra *et al.*, 2012). Specifically, a molecule-to-crystal difference in chemical shift is indicative of a combination of intermolecular interactions, notably hydrogen bonding and ring currents due to C–H··· π interactions, whereby the latter can be separately quantified by means of the nucleus independent chemical shift (NICS)

Table 3

Comparison of experimental ^1H chemical shifts with calculated^a (GIPAW) values (all in ppm) for the DI–PM cocrystal for the full crystal structure and an isolated dithianon or pyrimethanil molecule.

Atom	δ_{exp}	δ_{crystal}	δ_{molecule}	$\Delta\delta_{\text{crystal-molecule}}$
H1	7.4	7.4	7.8	−0.4
H2	6.2	5.9	7.4	−1.5
H3	7.7	7.6	7.4	0.2
H4	8.2	8.5	7.8	0.7
H17	9.1	9.7	9.2	0.5
H18	7.7	7.7	7.0	0.7
H19	7.8	7.3	6.6	0.7
H20	7.4	7.6	7.0	0.6
H21	8.0	8.4	6.4	2.0
H22/23/24 ^b	1.9	1.8	1.9	−0.1
H25	4.0	3.4	6.1	−2.7
H26/27/28 ^b	2.0	2.0	1.8	0.2
H29	9.1	10.5	6.9	3.6

Notes: (a) calculated isotropic chemical shieldings are determined from calculated chemical shieldings according to $\delta_{\text{calc}} = \sigma_{\text{ref}} - \sigma_{\text{calc}}$, where σ_{ref} equals 30.0 ppm; (b) for CH_3 groups, the calculated ^1H chemical shifts correspond to the average over the three H atoms.

(von Ragué Schleyer *et al.*, 1996; Sebastiani, 2006; Uldry *et al.*, 2008; Mafra *et al.*, 2012). Consider Table 3, which presents the change in ^1H chemical shift upon going from an isolated molecule to the full crystal, $\Delta\delta_{\text{crystal-molecule}}$, for the different H atoms in the DI–PM cocrystal. The largest positive change of 3.6 ppm is observed for the NH (H29) atom that is involved in an intermolecular $\text{N} \cdots \text{H} \cdots \text{O}$ hydrogen bond to atom O1 (see Fig. 1a; the $\text{N} \cdots \text{O}$ and $\text{H} \cdots \text{O}$ distances are 2.95 and 1.96 Å, respectively, with a 162° $\text{N} \cdots \text{H} \cdots \text{O}$ angle). Interestingly, $\Delta\delta_{\text{crystal-molecule}} = 2.0$ ppm for the aromatic CH H21 atom, for which Fig. 1(a) identifies an intermolecular $\text{C} \cdots \text{H} \cdots \text{O}$ so-called weak hydrogen-bonding (Desiraju & Steiner, 1999; Yates *et al.*, 2005; Uldry *et al.*, 2008) interaction (the $\text{C} \cdots \text{O}$ and $\text{H} \cdots \text{O}$ distances are 3.24 and 2.35 Å, respectively, with a 138° $\text{C} \cdots \text{H} \cdots \text{O}$ angle). The other H atoms, for which the magnitude of $\Delta\delta_{\text{crystal-molecule}}$ exceeds 1 ppm, are H25 (−2.7 ppm) and H2 (−1.6 ppm); as shown in Fig. 5, these marked changes

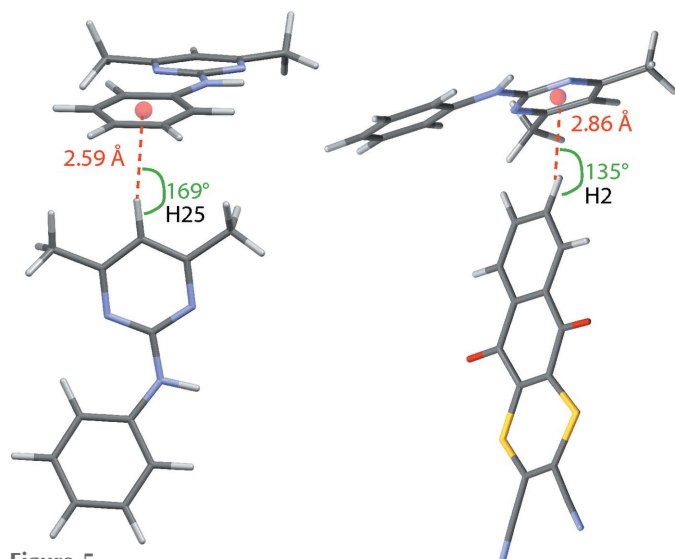


Figure 5

Schematic representations showing $\text{C} \cdots \text{H} \cdots \pi$ interactions for aromatic atoms (a) H25 and (b) H2.

Table 4

Comparison of calculated (GIPAW) NMR chemical shieldings (in ppm) for the DI–PM cocrystal for the full crystal structure and an isolated dithianon or pyrimethanil molecule.

Atom	σ_{molecule}	σ_{crystal}	$\sigma_{\text{crystal-molecule}}$
N1	−106.4	−88.9	17.5
N2	−107.2	−88.9	18.3
N9	98.9	91.4	−7.4
N10	−30.1	−33.0	−2.9
N11	−44.5	−42.4	2.1
O1	−363.4	−265.9	97.5
O2	−345.3	−322.2	23.1
S1	330.7	305.6	−25.1
S2	333.8	320.6	−13.2

in the ^1H chemical shift are a consequence of ring current effects associated with the proton pointing towards the centre of a six-membered aromatic ring of a nearby PM molecule in a $\text{C} \cdots \text{H} \cdots \pi$ interaction, as has been noted previously in a number of other cases (Brouwer *et al.*, 2008; Mafra *et al.*, 2012; Brown, 2012).

In the above discussion in §3.1, a close $\text{S} \cdots \text{O}$ distance, equal to 3.10 Å, between the O2 and S2 atoms of neighbouring DI molecules was noted; this is less than the sum of the van der Waals radii (3.32 Å) (Beno *et al.*, 2015; Zhang *et al.*, 2015). Indeed, there is a growing literature discussing $\text{S} \cdots \text{O}$ interactions (Burling & Goldstein, 1992; Iwaoka *et al.*, 2002; Beno *et al.*, 2015). While we have not carried out ^{17}O or ^{33}S solid-state NMR experiments as part of this study, an NMR crystallography approach enables the effect of such a putative $\text{S} \cdots \text{O}$ interaction on the oxygen and sulfur NMR chemical shieldings to be investigated by means of the GIPAW calculation that reports on all nuclei in the solid-state structure. An inspection of Table 4 shows that it is interesting that $\Delta\delta_{\text{crystal-molecule}}$ (note that this is the negative of the difference in calculated absolute shielding, with the latter being stated in Table 4) is much larger for O1 (−98 ppm), which is involved in a $\text{N} \cdots \text{H} \cdots \text{O}$ intermolecular hydrogen bonding, as compared to that for O2 (−23 ppm). Moreover, the change for S2 (13 ppm) is less than that for S1 (25 ppm), with both changes being small, though there is limited information on the range of experimentally observed solid-state NMR ^{33}S chemical shifts (Hansen *et al.*, 2008). We conclude that even though there is a close intermolecular $\text{S} \cdots \text{O}$ distance of 3.10 Å in the DI–PM cocrystal, there is not a marked effect on the calculated NMR chemical shieldings for the O2 and S2 nuclei.

4. Summary

In summary, we have presented here an NMR crystallography study of an agrochemical cocrystal. Specifically in combination with a GIPAW calculation of the NMR shieldings, ^1H – ^{13}C 2D correlation spectra enable the resolution and assignment of the NH, aromatic CH and methyl resonances for the DI–PM cocrystal, while specific intra- and intermolecular $\text{H} \cdots \text{H}$ proximities are identified in a ^1H DQ MAS spectrum. The performing of separate GIPAW calculations for the full crystal

structure and isolated DI and PM molecules yields the change in the NMR chemical shift upon going from the molecule to the crystal structure, thus allowing the quantitation of specific N—H···O, C—H···O and C—H··· π interactions.

Acknowledgements and funding information

ACP was supported by a Feodor Lynen Research Fellowship of the Alexander von Humboldt Foundation and a Newton International Fellowship of the Royal Society. EC and HP acknowledge funding from the Molecular Analytical Sciences Centre for Doctoral Training (EPSRC grant EP/L015307/1). We thank Peter Howe (Syngenta) for helpful discussions. Computational facilities were provided by the MidPlus Regional Centre of Excellence for Computational Science, Engineering and Mathematics, under EPSRC grant EP/K000128/1, and the University of Warwick Scientific Computing Research Technology Platform. The 700 MHz NMR spectrometer was partially funded from the European Research Council under the European Union's Seventh Framework Programme (FP/2007-2013)/ERC Grant Agreement 639907 (for Dr J. R. Lewandowski, Department of Chemistry, University of Warwick). The experimental and calculated data for this study are provided as a supporting data set from WRAP, the Warwick Research Archive Portal, at <http://wrap.warwick.ac.uk/85381>.

References

- Aakeroy, C. B. & Salmon, D. J. (2005). *CrystEngComm*, **7**, 439–448.
- Agilent (2014). *CrysAlis PRO*. Agilent Technologies Ltd, Yarnton, Oxfordshire, England.
- Ashbrook, S. E. & McKay, D. (2016). *Chem. Commun.* **52**, 7186–7204.
- Beno, B. R., Yeung, K. S., Bartberger, M. D., Pennington, L. D. & Meanwell, N. A. (2015). *J. Med. Chem.* **58**, 4383–4438.
- Betteridge, P. W., Carruthers, J. R., Cooper, R. I., Prout, K. & Watkin, D. J. (2003). *J. Appl. Cryst.* **36**, 1487.
- Bhatt, P. M., Azim, Y., Thakur, T. S. & Desiraju, G. R. (2009). *Cryst. Growth Des.* **9**, 951–957.
- Bielecki, A., Kolbert, A. C. & Levitt, M. H. (1989). *Chem. Phys. Lett.* **155**, 341–346.
- Blagden, N., de Matas, M., Gavan, P. T. & York, P. (2007). *Adv. Drug Deliv. Rev.* **59**, 617–630.
- Bonhomme, C., Gervais, C., Babonneau, F., Coelho, C., Pourpoint, F., Azais, T., Ashbrook, S. E., Griffin, J. M., Yates, J. R., Mauri, F. & Pickard, C. J. (2012). *Chem. Rev.* **112**, 5733–5779.
- Brouwer, D. H., Alavi, S. & Ripmeester, J. A. (2008). *Phys. Chem. Chem. Phys.* **10**, 3857–3860.
- Brown, S. P. (2007). *Prog. Nucl. Magn. Reson. Spectrosc.* **50**, 199–251.
- Brown, S. P. (2012). *Solid State Nucl. Magn. Reson.* **41**, 1–27.
- Brown, S. P., Zhu, X. X., Saalwachter, K. & Spiess, H. W. (2001). *J. Am. Chem. Soc.* **123**, 4275–4285.
- Burling, F. T. & Goldstein, B. M. (1992). *J. Am. Chem. Soc.* **114**, 2313–2320.
- Clark, S. J., Segall, M. D., Pickard, C. J., Hasnip, P. J., Probert, M. J., Refson, K. & Payne, M. C. (2005). *Z. Kristallogr.* **220**, 567–570.
- Cooper, R. I., Thompson, A. L. & Watkin, D. J. (2010). *J. Appl. Cryst.* **43**, 1100–1107.
- Desiraju, G. R. & Steiner, T. (1999). In *The Weak Hydrogen Bond in Structural Chemistry and Biology*. Oxford University Press.
- Dudenko, D. V., Yates, J. R., Harris, K. D. M. & Brown, S. P. (2013). *CrystEngComm*, **15**, 8797–8807.
- Elena, B., Pintacuda, G., Mifsud, N. & Emsley, L. (2006). *J. Am. Chem. Soc.* **128**, 9555–9560.
- Fung, B. M., Khitrin, A. K. & Ermolaev, K. (2000). *J. Magn. Reson.* **142**, 97–101.
- Hansen, M. R., Brorson, M., Bildsoe, H., Skibsted, J. & Jakobsen, H. J. (2008). *J. Magn. Reson.* **190**, 316–326.
- Harris, R. K. (2004). *Solid State Sci.* **6**, 1025–1037.
- Harris, R. K., Hodgkinson, P., Pickard, C. J., Yates, J. R. & Zorin, V. (2007). *Magn. Reson. Chem.* **45**, S174–S186.
- Harris, R. K., Wasylishen, R. E. & Duer, M. J. (2009). Editors. *NMR Crystallography*. Chichester: Wiley.
- Horst, J. H. ter, Deij, M. A. & Cains, P. W. (2009). *Cryst. Growth Des.* **9**, 1531–1537.
- Iwaoka, M., Takemoto, S. & Tomoda, S. (2002). *J. Am. Chem. Soc.* **124**, 10613–10620.
- Kerr, H. E., Softley, L. K., Suresh, K., Nangia, A., Hodgkinson, P. & Evans, I. R. (2015). *CrystEngComm*, **17**, 6707–6715.
- Lamberth, C., Jeanmart, S., Luksch, T. & Plant, A. (2013). *Science*, **341**, 742–746.
- Langer, B., Schnell, I., Spiess, H. W. & Grimmer, A. R. (1999). *J. Magn. Reson.* **138**, 182–186.
- Luedeker, D., Gossmann, R., Langer, K. & Brunklaus, G. (2016). *Cryst. Growth Des.* **16**, 3087–3100.
- Macrae, C. F., Edgington, P. R., McCabe, P., Pidcock, E., Shields, G. P., Taylor, R., Towler, M. & van de Streek, J. (2006). *J. Appl. Cryst.* **39**, 453–457.
- Mafra, L., Santos, S. M., Siegel, R., Alves, I., Paz, F. A. A., Dudenko, D. & Spiess, H. W. (2012). *J. Am. Chem. Soc.* **134**, 71–74.
- Metz, G., Wu, X. L. & Smith, S. O. (1994). *J. Magn. Reson. Ser. A*, **110**, 219–227.
- Palatinus, L. & Chapuis, G. (2007). *J. Appl. Cryst.* **40**, 786–790.
- Perdew, J. P., Burke, K. & Ernzerhof, M. (1996). *Phys. Rev. Lett.* **77**, 3865–3868.
- Pickard, C. J. & Mauri, F. (2001). *Phys. Rev. B* **63**, 245101.
- Pickard, C. J., Salager, E., Pintacuda, G., Elena, B. & Emsley, L. (2007). *J. Am. Chem. Soc.* **129**, 8932–8933.
- Ragué Schleyer, P. von, Maerker, C., Dransfeld, A., Jiao, H. & van Eikema Hommes, N. J. R. (1996). *J. Am. Chem. Soc.* **118**, 6317–6318.
- Sardo, M., Santos, S. M., Babaryk, A. A., Lopez, C., Alkorta, I., Elguero, J., Claramunt, R. M. & Mafra, L. (2015). *Solid State Nucl. Magn. Reson.* **65**, 49–63.
- Schmidt, J., Hoffmann, A., Spiess, H. W. & Sebastiani, D. (2006). *J. Phys. Chem. B*, **110**, 23204–23210.
- Schnell, I., Lupulescu, A., Hafner, S., Demco, D. E. & Spiess, H. W. (1998). *J. Magn. Reson.* **133**, 61–69.
- Sebastiani, D. (2006). *ChemPhysChem*, **7**, 164–175.
- Sommer, W., Gottwald, J., Demco, D. E. & Spiess, H. W. (1995). *J. Magn. Reson. Ser. A*, **113**, 131–134.
- Sowa, C., Saxell, H. E. & Vogel, R. (2013). EU Patent EP 2197278.
- Stevens, J. S., Byard, S. J., Seaton, C. C., Sadiq, G., Davey, R. J. & Schroeder, S. L. M. (2014). *Phys. Chem. Chem. Phys.* **16**, 1150–1160.
- Tatton, A. S., Pham, T. N., Vogt, F. G., Iuga, D., Edwards, A. J. & Brown, S. P. (2013). *Mol. Pharm.* **10**, 999–1007.
- Uldry, A. C., Griffin, J. M., Yates, J. R., Perez-Torrallba, M., Maria, M. D. S., Webber, A. L., Beaumont, M. L. L., Samoson, A., Claramunt, R. M., Pickard, C. J. & Brown, S. P. (2008). *J. Am. Chem. Soc.* **130**, 945–954.
- Vanderbilt, D. (1990). *Phys. Rev. B*, **41**, 7892.
- Watkin, D. J., Prout, C. K. & Pearce, L. J. (1996). *CAMERON*. Chemical Crystallography Laboratory, Oxford, England.
- Webber, A. L., Elena, B., Griffin, J. M., Yates, J. R., Pham, T. N., Mauri, F., Pickard, C. J., Gil, A. M., Stein, R., Lesage, A., Emsley, L. & Brown, S. P. (2010). *Phys. Chem. Chem. Phys.* **12**, 6970–6983.
- Webber, A. L., Emsley, L., Claramunt, R. M. & Brown, S. P. (2010). *J. Phys. Chem. A*, **114**, 10435–10442.
- Yates, J. R., Pham, T. N., Pickard, C. J., Mauri, F., Amado, A. M., Gil, A. M. & Brown, S. P. (2005). *J. Am. Chem. Soc.* **127**, 10216–10220.
- Yates, J. R., Pickard, C. J. & Mauri, F. (2007). *Phys. Rev. B*, **76**, 024401.
- Zhang, X., Gong, Z., Li, J. & Lu, T. (2015). *J. Chem. Inf. Model.* **55**, 2138–2153.

supporting information

Acta Cryst. (2017). C73, 149-156 [https://doi.org/10.1107/S2053229617000870]

Single-crystal X-ray diffraction and NMR crystallography of a 1:1 cocrystal of dithianon and pyrimethanil

Ann-Christin Pöppler, Emily K. Corlett, Harriet Pearce, Mark P. Seymour, Matthew Reid, Mark G. Montgomery and Steven P. Brown

Computing details

Data collection: *CrysAlis PRO* (Agilent, 2014); cell refinement: *CrysAlis PRO* (Agilent, 2014); data reduction: *CrysAlis PRO* (Agilent, 2014); program(s) used to solve structure: SUPERFLIP (Palatinus & Chapuis, 2007); program(s) used to refine structure: *CRYSTALS* (Betteridge *et al.*, 2003); molecular graphics: *CAMERON* (Watkin *et al.*, 1996) and *Mercury* (Macrae *et al.*, 2006); software used to prepare material for publication: *CRYSTALS* (Betteridge *et al.*, 2003).

5,10-Dioxo-5*H*,10*H*-naphtho[2,3-*b*][1,4]dithiine-2,3-dicarbonitrile-4,6-dimethyl-*N*-phenylpyrimidin-2-amine (1/1)

Crystal data

$C_{14}H_4N_2O_2S_2 \cdot C_{12}H_{13}N_3$
 $M_r = 495.59$
 Monoclinic, $P2_1/n$
 Hall symbol: -P 2yn
 $a = 7.1707$ (2) Å
 $b = 22.8006$ (6) Å
 $c = 13.8237$ (4) Å
 $\beta = 97.047$ (3)°
 $V = 2243.04$ (7) Å³
 $Z = 4$

$F(000) = 1024$
 $D_x = 1.467$ Mg m⁻³
 Cu $K\alpha$ radiation, $\lambda = 1.54184$ Å
 Cell parameters from 2711 reflections
 $\theta = 5.0\text{--}62.6^\circ$
 $\mu = 2.45$ mm⁻¹
 $T = 100$ K
 Plate, purple
 $0.60 \times 0.10 \times 0.02$ mm

Data collection

Agilent Xcalibur Onyx Ultra
 diffractometer
 Mirror monochromator
 $\omega/2\theta$ scans
 Absorption correction: multi-scan
 (CrysAlis PRO; Agilent, 2014)
 $T_{\min} = 0.596$, $T_{\max} = 1.000$
 5143 measured reflections

3160 independent reflections
 2667 reflections with $I > 2.0\sigma(I)$
 $R_{\text{int}} = 0.035$
 $\theta_{\max} = 58.9^\circ$, $\theta_{\min} = 3.2^\circ$
 $h = -5 \rightarrow 7$
 $k = -25 \rightarrow 25$
 $l = -14 \rightarrow 15$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.045$
 $wR(F^2) = 0.094$
 $S = 0.98$
 3141 reflections

109 parameters
 3 restraints
 Primary atom site location: other
 Hydrogen site location: difference Fourier map
 H atoms treated by a mixture of independent
 and constrained refinement

Method, part 1, Chebychev polynomial [weight]
 $= 1.0/[A_0*T_0(x) + A_1*T_1(x) \cdots + A_{n-1}*T_{n-1}(x)]$
 where A_i are the Chebychev coefficients listed
 below and $x = F/F_{\max}$ Method = Robust
 Weighting $W = [\text{weight}] * [1 - (\Delta F/6 * \sigma F)^2]^2$ A_i are: 0.138E + 04
 0.207E + 04 0.111E + 04 326.
 $(\Delta/\sigma)_{\max} = 0.001$
 $\Delta\rho_{\max} = 0.43 \text{ e } \text{\AA}^{-3}$
 $\Delta\rho_{\min} = -0.37 \text{ e } \text{\AA}^{-3}$

Special details

Experimental. The crystal was placed in the cold stream of an Oxford Cryosystems open-flow nitrogen cryostat (Cosier & Glazer, 1986) with a nominal stability of 0.1K.

Cosier, J. & Glazer, A.M., 1986. J. Appl. Cryst. 105-107.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (\AA^2)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
S1	0.57630 (11)	0.51819 (3)	0.79844 (6)	0.0223
C2	0.3685 (4)	0.48079 (13)	0.7529 (2)	0.0212
C3	0.2072 (4)	0.47646 (13)	0.7926 (2)	0.0202
S4	0.14929 (11)	0.50645 (3)	0.90297 (5)	0.0206
C5	0.3411 (4)	0.55384 (12)	0.9352 (2)	0.0177
C6	0.5006 (4)	0.55878 (12)	0.8951 (2)	0.0175
C7	0.6476 (4)	0.60247 (12)	0.9342 (2)	0.0179
O8	0.7809 (3)	0.61031 (9)	0.88850 (15)	0.0221
C9	0.6255 (4)	0.63243 (12)	1.0265 (2)	0.0167
C10	0.4604 (4)	0.62563 (12)	1.0704 (2)	0.0176
C11	0.3051 (4)	0.58862 (13)	1.0216 (2)	0.0176
O12	0.1533 (3)	0.58453 (9)	1.05284 (15)	0.0237
C13	0.4420 (4)	0.65186 (13)	1.1586 (2)	0.0214
C14	0.5889 (5)	0.68550 (14)	1.2043 (2)	0.0247
C15	0.7515 (5)	0.69313 (13)	1.1606 (2)	0.0235
C16	0.7717 (4)	0.66662 (13)	1.0719 (2)	0.0211
C17	0.0513 (5)	0.44419 (13)	0.7438 (2)	0.0227
N18	-0.0763 (4)	0.41868 (13)	0.7075 (2)	0.0334
C19	0.3871 (5)	0.45136 (14)	0.6633 (2)	0.0244
N20	0.4065 (4)	0.42677 (13)	0.5922 (2)	0.0372
N21	0.1269 (3)	0.65896 (11)	0.81861 (18)	0.0185
C22	0.1456 (4)	0.63048 (12)	0.7301 (2)	0.0193
C23	0.3046 (5)	0.63170 (13)	0.6818 (2)	0.0227
C24	0.3041 (5)	0.60183 (14)	0.5940 (2)	0.0281
C25	0.1478 (5)	0.57112 (14)	0.5532 (2)	0.0305
C26	-0.0103 (5)	0.56959 (14)	0.6017 (2)	0.0289
C27	-0.0132 (5)	0.59861 (13)	0.6889 (2)	0.0227
C28	0.2398 (4)	0.69992 (12)	0.8710 (2)	0.0172
N29	0.4071 (3)	0.71395 (10)	0.84429 (17)	0.0189
C30	0.5047 (4)	0.75525 (13)	0.8996 (2)	0.0204
C31	0.4355 (4)	0.78043 (13)	0.9782 (2)	0.0240

C32	0.2624 (5)	0.76203 (13)	1.0009 (2)	0.0233
N33	0.1612 (3)	0.72147 (11)	0.94755 (18)	0.0203
C34	0.1768 (5)	0.78633 (16)	1.0859 (3)	0.0350
C35	0.6919 (4)	0.77160 (14)	0.8698 (2)	0.0256
H131	0.3325	0.6473	1.1887	0.0272*
H141	0.5773	0.7037	1.2645	0.0304*
H151	0.8471	0.7168	1.1907	0.0275*
H161	0.8807	0.6722	1.0421	0.0261*
H231	0.4128	0.6519	0.7083	0.0274*
H241	0.4121	0.6020	0.5619	0.0339*
H251	0.1519	0.5519	0.4941	0.0370*
H261	−0.1170	0.5500	0.5748	0.0342*
H271	−0.1197	0.5972	0.7217	0.0260*
H311	0.5042	0.8085	1.0170	0.0294*
H342	0.0443	0.7782	1.0832	0.0544*
H341	0.1920	0.8277	1.0869	0.0549*
H343	0.2423	0.7710	1.1438	0.0548*
H352	0.7389	0.8070	0.9003	0.0411*
H353	0.6847	0.7771	0.8015	0.0418*
H351	0.7800	0.7420	0.8872	0.0415*
H211	0.022 (3)	0.6531 (11)	0.8421 (16)	0.0231*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
S1	0.0223	0.0220	0.0224	−0.0001	0.0015	−0.0055
C2	0.0274	0.0150	0.0197	0.0019	−0.0035	0.0014
C3	0.0236	0.0150	0.0203	−0.0003	−0.0043	0.0020
S4	0.0215	0.0185	0.0211	−0.0044	0.0004	−0.0018
C5	0.0202	0.0125	0.0189	0.0022	−0.0038	0.0041
C6	0.0214	0.0132	0.0169	0.0009	−0.0021	0.0022
C7	0.0197	0.0133	0.0204	0.0045	0.0009	0.0041
O8	0.0219	0.0229	0.0214	−0.0035	0.0027	−0.0011
C9	0.0188	0.0127	0.0171	0.0016	−0.0041	0.0010
C10	0.0199	0.0135	0.0184	0.0016	−0.0015	0.0037
C11	0.0187	0.0143	0.0198	0.0015	0.0023	0.0054
O12	0.0250	0.0224	0.0244	−0.0031	0.0062	0.0008
C13	0.0242	0.0199	0.0206	0.0015	0.0041	0.0008
C14	0.0338	0.0223	0.0175	−0.0007	0.0016	−0.0052
C15	0.0277	0.0184	0.0225	−0.0036	−0.0042	−0.0039
C16	0.0206	0.0174	0.0244	0.0004	−0.0003	0.0021
C17	0.0271	0.0205	0.0195	−0.0001	−0.0013	0.0002
N18	0.0413	0.0324	0.0255	−0.0073	0.0002	−0.0019
C19	0.0281	0.0203	0.0244	0.0027	0.0018	0.0001
N20	0.0423	0.0377	0.0314	0.0023	0.0037	−0.0076
N21	0.0157	0.0194	0.0204	−0.0024	0.0017	−0.0019
C22	0.0294	0.0114	0.0161	0.0038	−0.0013	0.0021
C23	0.0284	0.0181	0.0215	0.0000	0.0027	0.0018

C24	0.0429	0.0202	0.0223	0.0053	0.0090	0.0021
C25	0.0517	0.0213	0.0177	0.0021	0.0016	−0.0017
C26	0.0429	0.0167	0.0242	−0.0026	−0.0075	−0.0005
C27	0.0289	0.0183	0.0198	0.0015	−0.0020	0.0034
C28	0.0207	0.0113	0.0185	0.0042	−0.0016	0.0037
N29	0.0213	0.0152	0.0199	0.0012	0.0007	0.0033
C30	0.0215	0.0145	0.0238	0.0007	−0.0033	0.0047
C31	0.0283	0.0158	0.0267	−0.0028	−0.0013	−0.0036
C32	0.0270	0.0190	0.0235	0.0016	0.0019	−0.0027
N33	0.0224	0.0175	0.0210	0.0013	0.0029	−0.0031
C34	0.0366	0.0326	0.0373	−0.0031	0.0103	−0.0140
C35	0.0244	0.0239	0.0278	−0.0050	0.0004	0.0034

Geometric parameters (Å, °)

S1—C2	1.764 (3)	N21—H211	0.864 (17)
S1—C6	1.764 (3)	C22—C23	1.391 (4)
C2—C3	1.343 (4)	C22—C27	1.410 (4)
C2—C19	1.429 (4)	C23—C24	1.390 (4)
C3—S4	1.767 (3)	C23—H231	0.938
C3—C17	1.436 (4)	C24—C25	1.382 (5)
S4—C5	1.763 (3)	C24—H241	0.939
C5—C6	1.336 (4)	C25—C26	1.387 (5)
C5—C11	1.483 (4)	C25—H251	0.930
C6—C7	1.502 (4)	C26—C27	1.378 (4)
C7—O8	1.223 (4)	C26—H261	0.923
C7—C9	1.473 (4)	C27—H271	0.936
C9—C10	1.404 (4)	C28—N29	1.336 (4)
C9—C16	1.393 (4)	C28—N33	1.350 (4)
C10—C11	1.491 (4)	N29—C30	1.353 (4)
C10—C13	1.379 (4)	C30—C31	1.375 (4)
C11—O12	1.222 (4)	C30—C35	1.499 (4)
C13—C14	1.390 (4)	C31—C32	1.382 (5)
C13—H131	0.938	C31—H311	0.935
C14—C15	1.388 (5)	C32—N33	1.339 (4)
C14—H141	0.944	C32—C34	1.498 (4)
C15—C16	1.390 (4)	C34—H342	0.964
C15—H151	0.931	C34—H341	0.950
C16—H161	0.936	C34—H343	0.944
C17—N18	1.146 (4)	C35—H352	0.952
C19—N20	1.155 (4)	C35—H353	0.949
N21—C22	1.406 (4)	C35—H351	0.935
N21—C28	1.382 (4)		
C2—S1—C6	101.45 (14)	N21—C22—C27	115.5 (3)
S1—C2—C3	128.4 (2)	C23—C22—C27	119.0 (3)
S1—C2—C19	111.8 (2)	C22—C23—C24	119.6 (3)
C3—C2—C19	119.7 (3)	C22—C23—H231	120.6

C2—C3—S4	129.0 (2)	C24—C23—H231	119.7
C2—C3—C17	120.4 (3)	C23—C24—C25	121.3 (3)
S4—C3—C17	110.6 (2)	C23—C24—H241	119.7
C3—S4—C5	101.35 (14)	C25—C24—H241	118.9
S4—C5—C6	128.9 (2)	C24—C25—C26	119.0 (3)
S4—C5—C11	108.9 (2)	C24—C25—H251	119.1
C6—C5—C11	122.2 (3)	C26—C25—H251	121.9
S1—C6—C5	129.0 (2)	C25—C26—C27	120.8 (3)
S1—C6—C7	110.7 (2)	C25—C26—H261	120.2
C5—C6—C7	120.3 (3)	C27—C26—H261	118.9
C6—C7—O8	118.0 (3)	C22—C27—C26	120.2 (3)
C6—C7—C9	118.3 (3)	C22—C27—H271	119.3
O8—C7—C9	123.6 (3)	C26—C27—H271	120.5
C7—C9—C10	120.6 (3)	N21—C28—N29	120.3 (3)
C7—C9—C16	119.7 (3)	N21—C28—N33	112.5 (3)
C10—C9—C16	119.7 (3)	N29—C28—N33	127.2 (3)
C9—C10—C11	119.3 (3)	C28—N29—C30	115.5 (3)
C9—C10—C13	120.7 (3)	N29—C30—C31	121.7 (3)
C11—C10—C13	119.9 (3)	N29—C30—C35	115.9 (3)
C10—C11—C5	118.2 (3)	C31—C30—C35	122.4 (3)
C10—C11—O12	122.1 (3)	C30—C31—C32	118.3 (3)
C5—C11—O12	119.6 (3)	C30—C31—H311	121.5
C10—C13—C14	119.5 (3)	C32—C31—H311	120.2
C10—C13—H131	121.3	C31—C32—N33	121.7 (3)
C14—C13—H131	119.3	C31—C32—C34	122.1 (3)
C13—C14—C15	120.1 (3)	N33—C32—C34	116.2 (3)
C13—C14—H141	120.0	C28—N33—C32	115.6 (3)
C15—C14—H141	119.9	C32—C34—H342	113.2
C14—C15—C16	120.9 (3)	C32—C34—H341	108.8
C14—C15—H151	119.4	H342—C34—H341	107.6
C16—C15—H151	119.7	C32—C34—H343	108.7
C9—C16—C15	119.1 (3)	H342—C34—H343	110.3
C9—C16—H161	120.1	H341—C34—H343	108.1
C15—C16—H161	120.8	C30—C35—H352	111.7
C3—C17—N18	177.7 (3)	C30—C35—H353	111.4
C2—C19—N20	178.1 (3)	H352—C35—H353	107.6
C22—N21—C28	131.0 (3)	C30—C35—H351	110.5
C22—N21—H211	115.8 (12)	H352—C35—H351	107.8
C28—N21—H211	112.8 (12)	H353—C35—H351	107.7
N21—C22—C23	125.5 (3)		

Hydrogen-bond geometry (\AA , $^\circ$)

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
C23—H231 \cdots N29	0.94	2.36	2.950 (4)	121 (1)

C27—H271...O8 ⁱ	0.94	2.51	3.296 (4)	141 (1)
N21—H211...O8 ⁱ	0.86	2.15	2.985 (4)	162 (2)

Symmetry code: (i) $x-1, y, z$.